

# Produktinformation



Forschungsprodukte & Biochemikalien
Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

Weitere Information auf den folgenden Seiten! See the following pages for more information!



Lieferung & Zahlungsart siehe unsere Liefer- und Versandbedingungen

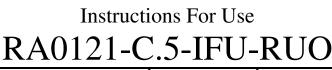
### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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**Revision: 1** 

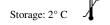
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P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.scytek.com

## GFAP (Astrocyte & Neural Stem Cell Marker); Clone GA-5 (Concentrate)

Availability/Contents:	<u>ltem #</u> BA0121-C.5	<u>Volume</u> 0.5 ml
Description:		
Species: Immunogen: Clone: Isotype: Entrez Gene ID: Hu Chromosome Loc.:	Mouse GFAP isolated from pig spinal cord GA-5 IgG1 2670 (Human); 14580 (Mouse); 24387 (Rat) 17q21.31	
Synonyms: Mol. Weight of Antigen: Format:	Astrocyte or Intermediate Filament Protein, Glial Fibrillary Acidic Protein (GFAP) ~50kDa 200μg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS	
Specificity:	with 0.05% BSA & 0.05% azide. This antibody recognizes a protein of ~50kDa which is identified as Glial Fibrillary Acidic Protein (GFAP). It shows no cross-reaction with other intermediate filament proteins.	
Background:	GFAP is specifically found in astroglia. GFAP is a very popular marker for localizing benign astrocyte and neoplastic cells of glial origin in the central nervous system. This antibody to GFAP is useful in differentiating primary gliomas from metastatic lesions in the brain and for documenting astrocytic differentiation in tumors outside the CNS.	
Species Reactivity: Positive Control: Cellular Localization: Titer/ Working Dilution:	Human, Mouse, Rat, Cow, Pig, Rabbit, Chicken. Others not known. Brain or Astrocytoma. Cytoplasmic Immunohistochemistry (Frozen and Formalin-fixed): 0.5-1 μg/ml Flow Cytometry: 0.5-1 μg/million cells Immunofluorescence: 1-2 μg/ml Western Blotting: 0.5-1 μg/ml	
Microbiological State:	Immunoprecipitation: This product is not sterile.	1-2 μg/500μg protein lysate









EC REP EmergoEurope (31)(0) 70 345-8570 Molsnstraat 15 2513 BH Hague, The Netherlands

Doc: IFU-Template2-8rev2



Ordering Information and Current Pricing at www.scytek.com

## Instructions For Use RA0121-C.5-IFU-RUO

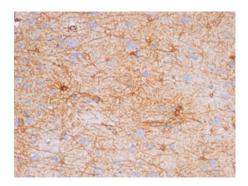
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**Uses/Limitations:** 

Not to be taken internally. For Research Use Only. This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin-fixed, paraffin-embedded tissue sections, to be viewed by light microscopy. Do not use if reagent becomes cloudy. Do not use past expiration date. Non-Sterile.



Formalin-paraffin human brain stained with GFAP; Clone GA-5. Note cytoplasmic staining.

#### Procedure:

- 1. **Tissue Section Pretreatment (Required):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500).
- Primary Antibody Incubation Time: We suggest an incubation period of 30 minutes at room temperature. However, depending upon the fixation conditions and the staining system employed, optimal incubation should be determined by the user.
- 3. **Visualization:** For maximum staining intensity we recommend the "UltraTek HRP Anti-Polyvalent Lab Pack" (ScyTek catalog# UHP125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).

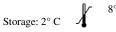
Precautions: Contains Sodium Azide as a preservative (0.09% w/v). Do not pipette by mouth. Avoid contact of reagents and specimens with skin and mucous membranes. Avoid microbial contamination of reagents or increased nonspecific staining may occur. This product contains no hazardous material at a <u>reportable concentration</u> according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.

#### **References:**

- 1. Herpers MJ et. Al. 1986, Acta Neuropathol, 70:333-339.
- 2. Van Muijen GN et. al. 1987, Lab Invest, 57:359-369.
- 3. Debus E, et. al. 1983, Differentiation, 25(2):193-203.

### Warranty:

No products or "Instructions For Use (IFU)" are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used.





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